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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/049,327	05/15/2002	Jay M Meythaler	UAB-15102/22	3596	
25006	7590 10/03/2003	20 10/03/2003		EXAMINER	
GIFFORD, KRASS, GROH, SPRINKLE			TRAVERS, RUSSELL S		
ANDERSON & CITKOWSKI, PC 280 N OLD WOODARD AVE			ART UNIT	PAPER NUMBER	
SUITE 400			1617	^	
BIRMINGHA	AM, MI 48009		DATE MAILED: 10/03/2003	9	

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary

Application No.

Applicant(s)

10/049,327

Examiner

R.S. Travers J.D., Ph.D.

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Meythaler et al



The MAILING DATE of this communication appea	ers on the cover sheet with the correspondence address			
Period for Reply				
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136 (a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the				
mailing date of this communication.				
 If the period for reply specified above is less than thirty (30) days, a reply withi If NO period for reply is specified above, the maximum statutory period will app 	by and will expire SIX (6) MONTHS from the mailing date of this communication.			
 Failure to reply within the set or extended period for reply will, by statute, caus Any reply received by the Office later than three months after the mailing date 				
earned patent term adjustment. See 37 CFR 1.704(b).	, , ,			
Status				
	ration is non-final			
	action is non-final.			
closed in accordance with the practice under Ex	e except for formal matters, prosecution as to the merits is parte Quayle, 1935 C.D. 11; 453 O.G. 213.			
Disposition of Claims				
4) 💢 Claim(s) <u>1-35</u>	is/are pending in the application.			
4a) Of the above, claim(s) 8-28	is/are withdrawn from consideration.			
5) Claim(s)	is/are allowed.			
6) X Claim(s) 1-7 and 29-35	is/are rejected.			
7) Claim(s)	is/are objected to.			
8) Claims	are subject to restriction and/or election requirement.			
Application Papers				
9) \square The specification is objected to by the Examiner.				
10) ☐ The drawing(s) filed on is/are a) ☐ accepted or b) ☐ objected to by the Examiner.				
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).				
) ☐ The proposed drawing correction filed on is: a) ☐ approved b) ☐ disapproved by the Examine				
If approved, corrected drawings are required in reply to this Office action.				
12) The oath or declaration is objected to by the Examiner.				
Priority under 35 U.S.C. §§ 119 and 120	· ·			
13) Acknowledgement is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).				
a) □ All b) □ Some* c) □ None of:				
1. Certified copies of the priority documents have been received.				
2. Certified copies of the priority documents have been received in Application No				
application from the International Bu				
*See the attached detailed Office action for a list of	the certified copies not received.			
14) Acknowledgement is made of a claim for domestic priority under 35 U.S.C. § 119(e).				
a) The translation of the foreign language provisional application has been received.				
15) ☐ Acknowledgement is made of a claim for domest	tic priority under 35 U.S.C. §§ 120 and/or 121.			
Attachment(s)				
1) X Notice of References Cited (PTO-892)	4) Interview Summary (PTO-413) Paper No(s).			
2) Notice of Draftsperson's Patent Drawing Review (PTO-948)	5) Notice of Informal Petent Application (PTO-152)			
3) X Information Disclosure Statement(s) (PTO-1449) Paper No(s). 4,5	6) Other:			

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The election filed September 4, 2003 has been received and entered into the file.

Claims 1-35 are presented for examination.

Applicant's election without traverse of Group I, claims 1-7 and 29-35 in Paper No. 8 is acknowledged.

Claims 8-28, reading on non-elected subject matter are withdrawn from consideration.

The following is a quotation of the first paragraph of 35 U.S.C. § 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

The specification is objected to under 35 U.S.C. § 112, first paragraph, as failing to adequately teach how to make and/or use the invention, and thereby failing to provide an enabling disclosure.

The instant specification fails to provide information that would allow the skilled artisan to practice the instant invention without undue experimentation. Attention is directed to *In re Wands*, 8 USPQ2d 1400 (CAFC 1988) at 1404 where the court set forth the eight factors to consider when assessing if a disclosure would have required undue experimentation. Citing *Ex parte Forman*, 230 USPQ 546 (BdApls 1986) at 547 the court recited eight factors:

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1) the quantity of experimentation necessary,

- 2) the amount of direction or guidance provided,
- 3) the presence of absence of working examples,
- 4) the nature of the invention,
- 5) the state of the prior art,
- 6) the relative skill of those in the art
- 7) the predictability of the art, and
- 8) the breadth of the claims.

Applicant fails to set forth the criteria that defines those "derivative" compounds which are useful for practicing the invention as claimed. Additionally, Applicant fails to provide information allowing the skilled artisan to ascertain these compounds without undue experimentation. In the instant case, only a limited number of "derivative" compound examples are set forth, thereby failing to provide sufficient working examples. It is noted that these examples are neither exhaustive, nor define the class of compounds required. The pharmaceutical art is unpredictable, requiring each embodiment to be individually assessed for physiological activity. The instant claims read on all "derivative" compounds which are useful for practicing the invention as claimed, necessitating an exhaustive search for the embodiments suitable to practice the claimed invention. Applicants fail to provide information sufficient to practice the claimed invention, absent undue experimentation.

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Claim 29 is rejected under 35 U.S.C. § 112, first paragraph, for the reasons set forth in the objection to the specification.

Claim 29 is rejected under 35 U.S.C. § 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 29 is rendered indefinite by the "derivative" and thereby failing to clearly set forth the metes and bounds of the patent protection desired. Criteria defining medicaments that are "derivative" compounds which are useful for practicing the invention as claimed are not set forth in the specification, thereby failing to provide information defining the instant inventions metes and bounds. Applicant's term fails to clearly define the subject matter encompassed by the instant claims, thus is properly rejected under 35 USC 112, second paragraph.

The specification is objected to under 35 U.S.C. § 112, first paragraph, as failing to adequately teach how to make and/or use the invention, and thereby failing to provide an enabling disclosure.

The instant specification fails to provide information that would allow the skilled artisan to practice the instant invention without undue experimentation. Attention is

directed to *In re Wands*, 8 USPQ2d 1400 (CAFC 1988) at 1404 where the court set forth the eight factors to consider when assessing if a disclosure would have required undue experimentation. Citing *Ex parte Forman*, 230 USPQ 546 (BdApls 1986) at 547 the court recited eight factors:

- 1) the quantity of experimentation necessary,
- 2) the amount of direction or guidance provided,
- 3) the presence of absence of working examples.
- 4) the nature of the invention,
- 5) the state of the prior art,
- 6) the relative skill of those in the art
- 7) the predictability of the art, and
- 8) the breadth of the claims.

Applicant fails to set forth the criteria that defines those maladies residing under the penumbra of "acquired disorders" envisioned as practiced in the invention as claimed. Additionally, Applicant fails to provide information allowing the skilled artisan to ascertain these compounds without undue experimentation. In the instant case, only a limited number of "acquired disorders" examples are set forth, thereby failing to provide sufficient working examples. It is noted that these examples are neither exhaustive, nor define the class of compounds required. The pharmaceutical art is unpredictable, requiring each embodiment to be individually assessed for physiological

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activity. The instant claims read on all "Acquired disorders", necessitating an exhaustive search for the embodiments suitable to practice the claimed invention.

Applicants fail to provide information sufficient to practice the claimed invention, absent undue experimentation.

Claim 35 is rejected under 35 U.S.C. § 112, first paragraph, for the reasons set forth in the objection to the specification.

Claim 35 is rejected under 35 U.S.C. § 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 35 is rendered indefinite by the phrase "acquired disorders" and thereby failing to clearly set forth the metes and bounds of the patent protection desired.

Criteria defining diseases residing under the penumbra of "acquired disorders" which are useful for practicing the invention as claimed are not set forth in the specification, thereby failing to provide information defining the instant inventions metes and bounds. Applicant's term fails to clearly define the subject matter encompassed by the instant claims, thus is properly rejected under 35 USC 112, second paragraph.

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The following is a quotation of the appropriate paragraphs of 35 U.S.C. § 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless --

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claims 1, 5-7, 29, 32, 33, 34 and 35 are rejected under 35 U.S.C. § 102(b) as being anticipated by Grilli et al.

The following is a quotation of 35 U.S.C. § 103 which forms the basis for all obviousness rejections set forth in this Office action:

A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

Subject matter developed by another person, which qualifies as prior art only under subsection (f) or (g) of section 102 of this title, shall not preclude patentability under this section where the subject matter and the claimed invention were, at the time the invention was made, owned by the same person or subject to an obligation of assignment to the same person.

Claims 1, 4-7, 29 and 31-35 are rejected under 35 U.S.C. § 103 as being unpatentable over Grilli et al, in view of the Merck Index.

Grilli et al teach the claimed non-steroidal anti-inflammatory compounds

(NSAID's) as old and well known in combination with various pharmaceutical carriers

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and excipients in a dosage form. These medicament are taught as useful for treating inflammation, and those maladies herein recited (see abstract and specification, especially pages 1-6) viewed by the skilled artisan as immuno-suppressive. Claims 4 and 31-32, and the primary reference, differ as to:

1) specific recitation of the claimed medicaments

This deficiency is cured by the Merck Index teaching Choline salicylate as an old and well known NSAID. Possessing the Grilli et la teaching of Parkinson's disease and Alzheimer's disease therapy with NSAID compounds, the skilled artisan would have been motivated to employ those NSAID compounds taught by the Merck Index for these conditions and enjoyed a reasonable expectation of therapeutic success, absent information to the contrary.

Claims 2, 3 and 30 are rejected under 35 U.S.C. § 103 as being unpatentable over Grilli et al, in view of the Merck Index, as set forth for claims 1, 4-7, 29 and 31-35, in further view of Jurna et al and Sakanashi et al.

Jurna et al and Sakanashi et al teach non-steroidal anti-inflammatory compounds (NSAID's) as old and well known in combination with various pharmaceutical carriers and excipients in a dosage form. These medicament are taught as useful for treating various maladies by administration via the interthecal and

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by inter-coronary injection. Claims 2, 3 and 30, and the primary references, differ as

to:

1) administration of these conventional medicaments by these routes

compounds by interthecal and inter-coronary injection. Jurna et al and Sakanashi et al

Claims 2, 3 and 30 specifically requires administration of the claimed NSAID

employed NSAID compounds by the interthecal and inter-coronary modes, not

specifically reciting another formulation, or mode of administration. The skilled artisan

would have seen interthecal and inter-coronary administration of NSAID compounds by

interthecal and inter-coronary routes as residing in the skilled artisan purview.

No claims are allowed.

Any inquiry concerning this communication should be directed to Russell

Travers at telephone number (703) 308-4603.

Russell Travers J.D., Ph.D.

Primary Examiner

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